White-Sutton syndrome

White-Sutton syndrome is a disorder that causes intellectual disability, specific facial features, and other signs and symptoms affecting various parts of the body. Most affected individuals have features of autism spectrum disorder (ASD), a varied condition characterized by impaired social skills, communication problems, and repetitive behaviors. However, in White-Sutton syndrome these features can occur along with other characteristics that are unusual in people with ASD, such as an overly friendly demeanor.

People with White-Sutton syndrome have delayed development, with speech and language usually being more delayed than motor skills such as walking. Intellectual disability can range from borderline normal to severe.

Most people with White-Sutton syndrome have mild abnormalities of the head and face, which can include an unusually small head (microcephaly); a wide, short skull (brachycephaly); wide-set eyes (hypertelorism); a flat or sunken appearance of the middle of the face (midface hypoplasia); and a small mouth with a thin upper lip.

A wide variety of additional signs and symptoms can occur with White-Sutton syndrome. Among the more common are hyperactivity; sleeping difficulties; vision defects, especially farsightedness; gastrointestinal problems; obesity; and short stature. Some individuals with White-Sutton syndrome are born with a hole in the muscle that separates the abdomen from the chest cavity (the diaphragm), which is called a diaphragmatic hernia.

Frequency

The prevalence of White-Sutton syndrome is unknown. Researchers estimate that changes in the gene associated with White-Sutton syndrome may account for up to 1 in 700 cases of intellectual disability, autism spectrum disorder, or both. However, most of these affected individuals have not been diagnosed with White-Sutton syndrome and may not exhibit all the features of this disorder.

Causes

White-Sutton syndrome is caused by mutations in the *POGZ* gene. This gene provides instructions for making a protein that is found in the cell nucleus. The POGZ protein attaches (binds) to chromatin, which is the network of DNA and proteins that packages DNA into chromosomes. Binding of the POGZ protein is part of the process that changes the structure of chromatin (chromatin remodeling) to alter how tightly regions of DNA are packaged. Chromatin remodeling is one way gene activity (expression) is regulated; when DNA is tightly packed gene expression is lower than when DNA is loosely packed. Regulation of gene expression by the POGZ protein is thought to be

important to brain development, but the specific function of POGZ in the brain is not well understood.

POGZ gene mutations are thought to impair the ability of the POGZ protein to bind to chromatin, leading to abnormal gene expression that affects development of the brain and other body systems. However, little is known about the specific changes in gene expression and how they lead to the development of intellectual disability and other signs and symptoms of White-Sutton syndrome.

Inheritance Pattern

This condition is inherited in an autosomal dominant pattern, which means one copy of the altered gene in each cell is sufficient to cause the disorder.

Most cases of this condition occur in people with no history of the disorder in their family, and result from new (de novo) mutations in the gene that occur during the formation of reproductive cells (eggs or sperm) or in early embryonic development.

Other Names for This Condition

- mental retardation, autosomal dominant 37
- MRD37
- WHSUS

Diagnosis & Management

Genetic Testing Information

- What is genetic testing?
 /primer/testing/genetictesting
- Genetic Testing Registry: White-sutton syndrome https://www.ncbi.nlm.nih.gov/qtr/conditions/C4225351/

Other Diagnosis and Management Resources

- Agency for Healthcare Research and Quality: Therapies for Children with Autism Spectrum Disorder https://effectivehealthcare.ahrq.gov/products/autism-update/consumer/
- Eunice Kennedy Shriver National Institute of Child Health and Human Development: How Do Health Care Providers Diagnose IDDs? https://www.nichd.nih.gov/health/topics/idds/conditioninfo/diagnosed
- Eunice Kennedy Shriver National Institute of Child Health and Human Development: What Are Treatments for IDDs? https://www.nichd.nih.gov/health/topics/idds/conditioninfo/cure

Additional Information & Resources

Health Information from MedlinePlus

- Encyclopedia: Autism Spectrum Disorder https://medlineplus.gov/ency/article/001526.htm
- Encyclopedia: Intellectual Disability https://medlineplus.gov/ency/article/001523.htm
- Health Topic: Autism Spectrum Disorder https://medlineplus.gov/autismspectrumdisorder.html
- Health Topic: Developmental Disabilities https://medlineplus.gov/developmentaldisabilities.html

Additional NIH Resources

- Eunice Kennedy Shriver National Institute of Child Health and Human Development: How Do Health Care Providers Diagnose IDDs? https://www.nichd.nih.gov/health/topics/idds/conditioninfo/diagnosed
- Eunice Kennedy Shriver National Institute of Child Health and Human Development: What Are Treatments for IDDs? https://www.nichd.nih.gov/health/topics/idds/conditioninfo/cure
- National Human Genome Research Institute: Learning About Autism https://www.genome.gov/Genetic-Disorders/Autism
- National Institute of Mental Health: Autism Spectrum Disorder https://www.nimh.nih.gov/health/topics/autism-spectrum-disorders-asd/index.shtml
- National Institute of Neurological Disorders and Stroke: Autism Spectrum Disorder Fact Sheet https://www.ninds.nih.gov/Disorders/Patient-Caregiver-Education/Fact-Sheets/ Autism-Spectrum-Disorder-Fact-Sheet

Educational Resources

- American Academy of Child and Adolescent Psychiatry: Intellectual Disabilities https://www.aacap.org/AACAP/Families_and_Youth/Facts_for_Families/FFF-Guide/Children-with-an-Intellectual-Disability-023.aspx
- American Academy of Family Physicians: Caring for a Person Who Has Intellectual or Developmental Disabilities https://familydoctor.org/caring-for-a-person-who-has-intellectual-or-developmentaldisabilities/?adfree=true
- Centers for Disease Control and Prevention: Autism Spectrum Disorder https://www.cdc.gov/ncbddd/autism/index.html
- Centers for Disease Control and Prevention: Facts About Intellectual Disability https://www.cdc.gov/ncbddd/actearly/pdf/parents_pdfs/IntellectualDisability.pdf

- MalaCards: white-sutton syndrome https://www.malacards.org/card/white_sutton_syndrome
- TeensHealth: What is Autism? https://kidshealth.org/en/teens/autism.html

Patient Support and Advocacy Resources

- American Association on Intellectual and Developmental Disabilities (AAIDD) https://www.aaidd.org/
- Autism Society https://www.autism-society.org/
- Autism Speaks https://www.autismspeaks.org/
- White Sutton Syndrome Foundation https://whitesutton.org/

Scientific Articles on PubMed

PubMed
 https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28pogz%5BTIAB%5D
 %29+AND+%28intellectual+disability%5BTIAB%5D%29%29

Catalog of Genes and Diseases from OMIM

 WHITE-SUTTON SYNDROME http://omim.org/entry/616364

Medical Genetics Database from MedGen

 White-sutton syndrome https://www.ncbi.nlm.nih.gov/medgen/897984

Sources for This Summary

- Dentici ML, Niceta M, Pantaleoni F, Barresi S, Bencivenga P, Dallapiccola B, Digilio MC, Tartaglia M. Expanding the phenotypic spectrum of truncating POGZ mutations: Association with CNS malformations, skeletal abnormalities, and distinctive facial dysmorphism. Am J Med Genet A. 2017 May 7. doi: 10.1002/ajmg.a.38255. [Epub ahead of print]
 Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/28480548
- Stessman HA, Willemsen MH, Fenckova M, Penn O, Hoischen A, Xiong B, Wang T, Hoekzema K, Vives L, Vogel I, Brunner HG, van der Burgt I, Ockeloen CW, Schuurs-Hoeijmakers JH, Klein Wassink-Ruiter JS, Stumpel C, Stevens SJ, Vles HS, Marcelis CM, van Bokhoven H, Cantagrel V, Colleaux L, Nicouleau M, Lyonnet S, Bernier RA, Gerdts J, Coe BP, Romano C, Alberti A, Grillo L, Scuderi C, Nordenskjöld M, Kvarnung M, Guo H, Xia K, Piton A, Gerard B, Genevieve D, Delobel B, Lehalle D, Perrin L, Prieur F, Thevenon J, Gecz J, Shaw M, Pfundt R, Keren B, Jacquette A, Schenck A, Eichler EE, Kleefstra T. Disruption of POGZ Is Associated with Intellectual Disability and Autism Spectrum Disorders. Am J Hum Genet. 2016 Mar 3;98(3):541-52. doi: 10.1016/j.ajhg.2016.02.004.

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Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4890241/

- Tan B, Zou Y, Zhang Y, Zhang R, Ou J, Shen Y, Zhao J, Luo X, Guo J, Zeng L, Hu Y, Zheng Y, Pan Q, Liang D, Wu L. A novel de novo POGZ mutation in a patient with intellectual disability. J Hum Genet. 2016 Apr;61(4):357-9. doi: 10.1038/jhg.2015.156. Epub 2016 Jan 14. Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/26763879
- White J, Beck CR, Harel T, Posey JE, Jhangiani SN, Tang S, Farwell KD, Powis Z, Mendelsohn NJ, Baker JA, Pollack L, Mason KJ, Wierenga KJ, Arrington DK, Hall M, Psychogios A, Fairbrother L, Walkiewicz M, Person RE, Niu Z, Zhang J, Rosenfeld JA, Muzny DM, Eng C, Beaudet AL, Lupski JR, Boerwinkle E, Gibbs RA, Yang Y, Xia F, Sutton VR. POGZ truncating alleles cause syndromic intellectual disability. Genome Med. 2016 Jan 6;8(1):3. doi: 10.1186/s13073-015-0253-0. Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/26739615
 Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4702300/
- Ye Y, Cho MT, Retterer K, Alexander N, Ben-Omran T, Al-Mureikhi M, Cristian I, Wheeler PG, Crain C, Zand D, Weinstein V, Vernon HJ, McClellan R, Krishnamurthy V, Vitazka P, Millan F, Chung WK. De novo POGZ mutations are associated with neurodevelopmental disorders and microcephaly. Cold Spring Harb Mol Case Stud. 2015 Oct;1(1):a000455. doi: 10.1101/mcs.a000455. Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/27148570 Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4850885/

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